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NEWS 6 MAY 19 Derwent World Patents Index to be reloaded and enhanced  
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NEWS 8 MAY 30 The F-Term thesaurus is now available in CA/CAPLUS  
NEWS 9 JUN 02 The first reclassification of IPC codes now complete in  
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NEWS 10 JUN 26 TULSA/TULSA2 reloaded and enhanced with new search and  
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NEWS 13 JUL 14 FSTA enhanced with Japanese patents  
NEWS 14 JUL 19 Coverage of Research Disclosure reinstated in DWPI  
NEWS 15 AUG 09 INSPEC enhanced with 1898-1968 archive  
NEWS 16 AUG 28 ADISCTI Reloaded and Enhanced  
NEWS 17 AUG 30 CA(SM)/CAPLUS(SM) Austrian patent law changes  
NEWS 18 SEP 11 CA/CAPLUS enhanced with more pre-1907 records  
NEWS 19 SEP 21 CA/CAPLUS fields enhanced with simultaneous left and right  
truncation  
NEWS 20 SEP 25 CA(SM)/CAPLUS(SM) display of CA Lexicon enhanced  
NEWS 21 SEP 25 CAS REGISTRY(SM) no longer includes Concord 3D coordinates  
NEWS 22 SEP 25 CAS REGISTRY(SM) updated with amino acid codes for pyrrolysine  
  
NEWS EXPRESS JUNE 30 CURRENT WINDOWS VERSION IS V8.01b, CURRENT  
MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),  
AND CURRENT DISCOVER FILE IS DATED 26 JUNE 2006.  
  
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FILE 'HOME' ENTERED AT 10:33:59 ON 26 SEP 2006

=> FIL HCAPLUS

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
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FULL ESTIMATED COST

FILE 'HCAPLUS' ENTERED AT 10:34:25 ON 26 SEP 2006

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FILE COVERS 1907 - 26 Sep 2006 VOL 145 ISS 14

FILE LAST UPDATED: 25 Sep 2006 (20060925/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s diaryl carbonate

14380 DIARYL

92 DIARYLS

14435 DIARYL

(DIARYL OR DIARYLS)

286399 CARBONATE

66944 CARBONATES

319101 CARBONATE

(CARBONATE OR CARBONATES)

L1 711 DIARYL CARBONATE

(DIARYL(W) CARBONATE)

=> s l1 and process

2312002 PROCESS

1569204 PROCESSES

3450734 PROCESS

(PROCESS OR PROCESSES)

L2 197 L1 AND PROCESS

=> s l2 and tempering

25442 TEMPERING

55 TEMPERINGS

25458 TEMPERING

(TEMPERING OR TEMPERINGS)

09/26/2006 10791587.trn

L3 0 L2 AND TEMPERING

=> s l1 and tempering  
25442 TEMPERING  
55 TEMPERINGS  
25458 TEMPERING  
(TEMPERING OR TEMPERINGS)

L4 1 L1 AND TEMPERING

=> s l1 and thermal pre-treatment  
1062396 THERMAL  
72 THERMALS  
1062429 THERMAL  
(THERMAL OR THERMALS)  
205129 PRE  
691 PRES  
205491 PRE  
(PRE OR PRES)  
2183793 TREATMENT  
203094 TREATMENTS  
2291777 TREATMENT  
(TREATMENT OR TREATMENTS)  
144 THERMAL PRE-TREATMENT  
(THERMAL(W) PRE(W) TREATMENT)  
L5 0 L1 AND THERMAL PRE-TREATMENT

=> s l2 and thermal pre-treatment  
1062396 THERMAL  
72 THERMALS  
1062429 THERMAL  
(THERMAL OR THERMALS)  
205129 PRE  
691 PRES  
205491 PRE  
(PRE OR PRES)  
2183793 TREATMENT  
203094 TREATMENTS  
2291777 TREATMENT  
(TREATMENT OR TREATMENTS)  
144 THERMAL PRE-TREATMENT  
(THERMAL(W) PRE(W) TREATMENT)  
L6 0 L2 AND THERMAL PRE-TREATMENT

=> s l2 and thermal treatment  
1062396 THERMAL  
72 THERMALS  
1062429 THERMAL  
(THERMAL OR THERMALS)  
2183793 TREATMENT  
203094 TREATMENTS  
2291777 TREATMENT  
(TREATMENT OR TREATMENTS)  
37525 THERMAL TREATMENT  
(THERMAL(W) TREATMENT)  
L7 0 L2 AND THERMAL TREATMENT

=> s l1 and thermal treatment  
1062396 THERMAL  
72 THERMALS

09/26/2006 10791587.trn

1062429 THERMAL  
(THERMAL OR THERMALS)  
2183793 TREATMENT  
203094 TREATMENTS  
2291777 TREATMENT  
(TREATMENT OR TREATMENTS)  
37525 THERMAL TREATMENT  
(THERMAL(W) TREATMENT)

L8 0 L1 AND THERMAL TREATMENT

=> s l1 and thermal

1062396 THERMAL  
72 THERMALS  
1062429 THERMAL  
(THERMAL OR THERMALS)

L9 38 L1 AND THERMAL

=> s l2 and thermal

1062396 THERMAL  
72 THERMALS  
1062429 THERMAL  
(THERMAL OR THERMALS)

L10 5 L2 AND THERMAL

=> d his

(FILE 'HOME' ENTERED AT 10:33:59 ON 26 SEP 2006)

FILE 'HCAPLUS' ENTERED AT 10:34:25 ON 26 SEP 2006

L1 711 S DIARYL CARBONATE  
L2 197 S L1 AND PROCESS  
L3 0 S L2 AND TEMPERING  
L4 1 S L1 AND TEMPERING  
L5 0 S L1 AND THERMAL PRE-TREATMENT  
L6 0 S L2 AND THERMAL PRE-TREATMENT  
L7 0 S L2 AND THERMAL TREATMENT  
L8 0 S L1 AND THERMAL TREATMENT  
L9 38 S L1 AND THERMAL  
L10 5 S L2 AND THERMAL

=> s l2 and catalyst system

735799 CATALYST  
738463 CATALYSTS  
944541 CATALYST  
(CATALYST OR CATALYSTS)

2325088 SYSTEM  
1273386 SYSTEMS  
3151860 SYSTEM  
(SYSTEM OR SYSTEMS)

16897 CATALYST SYSTEM  
(CATALYST(W) SYSTEM)

L11 17 L2 AND CATALYST SYSTEM

=> s l2 and catalyst

735799 CATALYST  
738463 CATALYSTS  
944541 CATALYST  
(CATALYST OR CATALYSTS)

L12 124 L2 AND CATALYST

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L4 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2006 ACS on STN  
ACCESSION NUMBER: 1993:125243 HCAPLUS  
DOCUMENT NUMBER: 118:125243  
TITLE: Preparation of cyclic carbonate esters  
INVENTOR(S): Schoen, Norbert; Buysch, Hans Josef; Leitz, Edgar;  
Ott, Karl Heinz  
PATENT ASSIGNEE(S): Bayer A.-G., Germany  
SOURCE: Ger. Offen., 7 pp.  
CODEN: GWXXBX  
DOCUMENT TYPE: Patent  
LANGUAGE: German  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 4109236	A1	19920924	DE 1991-4109236	19910321
PRIORITY APPLN. INFO.:			DE 1991-4109236	19910321
OTHER SOURCE(S):	MARPAT 118:125243			

AB Cyclic carbonates (d.p. 1 or 2) are prepared simply, in satisfactory quality for anionic polymerization, by catalyzed transesterification of C3-18 diols with diaryl carbonates, tempering, and depolymn.  
Heating 14.0 mol neopentyl glycol, 14.0 mol (PhO)2CO, and 730 mg Bu2Sn dilaurate at 110-190°/30-20 mbar with distillation of PhOH, heating the pot residue at 195-210°/300-450 mbar in a stream of N for 18 h, and distillation in vacuo gave 87% cyclic carbonate (I) with purity 99.5%. Anionic polymerization of I gave a polycarbonate in 96% yield with mol. weight 35,000; vs. 70 and 5000, resp., when I (purity 95%) was prepared without the tempering step.

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L10 ANSWER 1 OF 5 HCAPLUS COPYRIGHT 2006 ACS on STN  
ACCESSION NUMBER: 2004:756014 HCAPLUS  
DOCUMENT NUMBER: 141:279423  
TITLE: Carbonylation process and catalysts for the production of a diaryl carbonates from phenols  
INVENTOR(S): Dahlmann, Marg; Fischer, Peter; Hansen, Sven-Michael; Reisinger, Claus-Peter  
PATENT ASSIGNEE(S): Bayer Materialscience AG, Germany  
SOURCE: Ger. Offen., 8 pp.  
CODEN: GWXXBX  
DOCUMENT TYPE: Patent  
LANGUAGE: German  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 10309954	A1	20040916	DE 2003-10309954	20030307
EP 1460055	A1	20040922	EP 2004-4639	20040301
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,				

IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK  
 US 2004192953 A1 20040930 US 2004-791587 20040302  
 CN 1526694 A 20040908 CN 2004-10008006 20040305  
 JP 2004269530 A2 20040930 JP 2004-61940 20040305  
 PRIORITY APPLN. INFO.: DE 2003-10309954 A. 20030307

OTHER SOURCE(S): CASREACT 141:279423; MARPAT 141:279423

AB A procedure is described for the production of diaryl carbonates (e.g., di-Ph carbonate) by the direct carbonylation of phenols (e.g., phenol) in the presence of a catalyst system where the catalyst system is activated by thermal pretreatment in a sep. reaction apparatus

L10 ANSWER 2 OF 5 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2001:886302 HCAPLUS

DOCUMENT NUMBER: 136:20376

TITLE: Wholly aromatic polyester-polycarbonates and their production process

INVENTOR(S): Sakurai, Hiroshi; Ishiwata, Toyoaki; Miyoshi, Takanori; Matsumura, Shunichi

PATENT ASSIGNEE(S): Teijin Limited, Japan

SOURCE: PCT Int. Appl., 26 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 3

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001092370	A1	20011206	WO 2001-JP4611	20010531
W: CA, CN, ID, JP, KR, SG, US				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR				
EP 1291374	A1	20030312	EP 2001-934464	20010531
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI, CY, TR				
US 2003181627	A1	20030925	US 2002-297082	20021202
US 6858701	B2	20050222		
PRIORITY APPLN. INFO.:			JP 2000-164532	A 20000601
			JP 2000-260581	A 20000830
			JP 2001-1333	A 20010109
			WO 2001-JP4611	W 20010531

OTHER SOURCE(S): MARPAT 136:20376

AB The polymers have a satisfactory color tone, excellent thermal stability, and an alkali metal content of  $\leq 10$  ppm. The polymers are advantageously produced by reacting aromatic dicarboxylic acids (e.g., terephthalic acid), aromatic diols (e.g., bisphenol A), and diaryl carbonates (e.g., di-Ph carbonate) in a specific molar proportion using a pyridine compound (e.g., 4-dimethylaminopyridine) as a catalyst.

REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L10 ANSWER 3 OF 5 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1995:276802 HCAPLUS

DOCUMENT NUMBER: 122:32363

TITLE: Process for manufacture of thermoplastic polycarbonates

INVENTOR(S): Kauth, Hermann; Kuehling, Steffen; Alewelt, Wolfgang; Freitag, Dieter

PATENT ASSIGNEE(S): Bayer A.-G., Germany  
 SOURCE: Ger. Offen., 7 pp.  
 CODEN: GWXXBX  
 DOCUMENT TYPE: Patent  
 LANGUAGE: German  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 4240314	A1	19940609	DE 1992-4240314	19921201
US 5373082	A	19941213	US 1993-155309	19931119
JP 06220184	A2	19940809	JP 1993-317499	19931125
JP 3174446	B2	20010611		
BE 1007762	A3	19951017	BE 1993-1304	19931125
NL 9302064	A	19940701	NL 1993-2064	19931129
NL 194866	B	20030106		
NL 194866	C	20030506		

PRIORITY APPLN. INFO.: DE 1992-4240314 A 19921201  
 AB In polycarbonate manufacture by melt transesterification, polycarbonates (especially

waste polycarbonates) from aromatic bisphenols are dissolved in monophenols; degraded to oligocarbonates, diaryl carbonates, and bisphenols in the presence of quaternary ammonium or phosphonium catalysts at 100-295°. After optional removal of fillers and/or additives, high-viscosity oligocarbonates (mol. weight 8000-18,000) are prepared and polycondensed at 250-295° and 0.1-500 mbar to polycarbonates with weight-average mol. weight 20,000-100,000. Thus, a bisphenol A polycarbonate

was

dissolved in PhOH and degraded at 180° in the presence of Me4NOH to an oligocarbonate, which was then polycondensed at 280° to a solvent-free polycarbonate with solution viscosity 1.269 (CH2Cl2, 5 g/L, 25°) and branch unit content 15 ppm.

L10 ANSWER 4 OF 5 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1993:650161 HCAPLUS

DOCUMENT NUMBER: 119:250161

TITLE: Process for making a silylisocyanurate from the catalytic thermal trimerization of silylorganocarbamate

INVENTOR(S): Pepe, Enrico J.; Su, Shiu Chin H.; Turner, Scot M.

PATENT ASSIGNEE(S): Union Carbide Chemicals and Plastics Technology Corp., USA

SOURCE: U.S., 9 pp.  
 CODEN: USXXAM

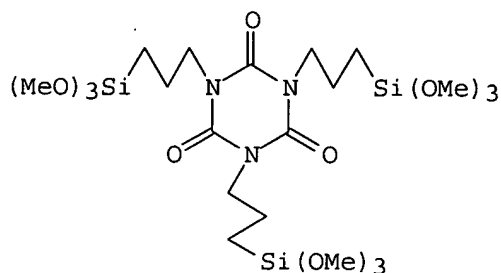
DOCUMENT TYPE: Patent  
 LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5218133	A	19930608	US 1992-932584	19920820
EP 583581	A1	19940223	EP 1993-110257	19930628
EP 583581	B1	19970528		
R: DE, FR, GB, IT, NL				
JP 06228166	A2	19940816	JP 1993-178476	19930628
JP 2963309	B2	19991018		
JP 10067788	A2	19980310	JP 1997-195226	19970707

JP 2916442 B2 19990705  
 PRIORITY APPLN. INFO.: US 1992-932584 A 19920820  
 JP 1993-178476 A3 19930628  
 OTHER SOURCE(S): MARPAT 119:250161  
 GI



AB A process for making a silylorganocarbamate or a silylisocyanurate comprises reacting an aminosilane with a dialkyl carbonate, diaryl carbonate or a mixture thereof in the presence of a basic catalyst to obtain the silylorganocarbamate; optionally, neutralizing the basic catalyst and residual aminosilane with a neutralizing agent; and adding a cracking catalyst and heating at subatmospheric pressure to obtain the silylisocyanurate e.g., I, or heating a silylorganocarbamate at a temperature sufficient for dissociation of the carbamate at subatmospheric pressure in the presence of a cracking catalyst and a trimerization catalyst to obtain a silylisocyanurate.

L10 ANSWER 5 OF 5 HCAPLUS COPYRIGHT 2006 ACS on STN  
 ACCESSION NUMBER: 1992:60254 HCAPLUS  
 DOCUMENT NUMBER: 116:60254  
 TITLE: Aromatic polycarbonates with controlled molecular weight and hydroxy end group content  
 INVENTOR(S): Fukawa, Isaburo; Tanabe, Tsuneaki  
 PATENT ASSIGNEE(S): Asahi Chemical Industry Co., Ltd., Japan  
 SOURCE: Jpn. Kokai Tokkyo Koho, 17 pp.  
 CODEN: JKXXAF  
 DOCUMENT TYPE: Patent  
 LANGUAGE: Japanese  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 03252421	A2	19911111	JP 1990-47327	19900301
JP 07098862	B4	19951025		

PRIORITY APPLN. INFO.: JP 1990-47327 19900301

AB The title polymers are prepared by thermal prepolymerization of dihydroxydiaryl compounds with diaryl carbonates, determination of the mol. weight of the polymers and the end group composition, and adding the dihydroxydiaryl compound and/or the diaryl carbonate to obtain the desired mol. weight and content of OH end groups. The process is demonstrated by polymerizing bisphenol A with di-Ph carbonate.



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L11 ANSWER 1 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2005:349044 HCAPLUS

DOCUMENT NUMBER: 142:394138

TITLE: Water-resistant carbonylation catalyst system for the production of diaryl carbonates via the direct carbonylation of phenolic compounds

INVENTOR(S): Soloveichik, Grigorii Lev; Chuck, Timothy Leigh; Shalyaev, Kirill Vladimirovich; Pressman, Eric James; Bonitatebus, Peter John

PATENT ASSIGNEE(S): General Electric Company, USA

SOURCE: U.S. Pat. Appl. Publ., 9 pp.

CODEN: USXXCO

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2005085656	A1	20050421	US 2003-687411	20031015
US 7084291	B2	20060801		
WO 2005040089	A2	20050506	WO 2004-US30610	20040917

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.: US 2003-687411 A 20031015

OTHER SOURCE(S): CASREACT 142:394138

AB A method of increasing the amount of diaryl carbonates (e.g., di-Ph carbonate) produced per amount of catalyst consumed in a phenolic compound (e.g., phenol) carbonylation process is described. Phenolic compound carbonylation produces water as a reaction byproduct which reduces the turnover number (TON) of the catalyst. A mixture of a phenolic precursor, a base-containing catalyst and co-catalyst components and at least one chemical additive comprising a halide or hydroxide of alkali metal or alkaline earth metal when carbonylated together under specific conditions increases the TON and water resistivity of a palladium catalyst. The metal halide likely makes the catalyst less susceptible to degradation by water hence increasing the reaction yield per weight of catalyst consumed.

REFERENCE COUNT: 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 2 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2004:756014 HCAPLUS

DOCUMENT NUMBER: 141:279423

TITLE: Carbonylation process and catalysts for the

09/26/2006

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production of a diaryl carbonates  
from phenols

INVENTOR(S): Dahlmann, Marc; Fischer, Peter; Hansen, Sven-Michael;  
Reisinger, Claus-Peter  
PATENT ASSIGNEE(S): Bayer Materialscience Ag, Germany  
SOURCE: Ger. Offen., 8 pp.  
CODEN: GWXXBX  
DOCUMENT TYPE: Patent  
LANGUAGE: German  
FAMILY ACC. NUM. COUNT: 1  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 10309954	A1	20040916	DE 2003-10309954	20030307
EP 1460055	A1	20040922	EP 2004-4639	20040301
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK				
US 2004192953	A1	20040930	US 2004-791587	20040302
CN 1526694	A	20040908	CN 2004-10008006	20040305
JP 2004269530	A2	20040930	JP 2004-61940	20040305
PRIORITY APPLN. INFO.:			DE 2003-10309954	A 20030307

OTHER SOURCE(S): CASREACT 141:279423; MARPAT 141:279423

AB A procedure is described for the production of diaryl carbonates (e.g., di-Ph carbonate) by the direct carbonylation of phenols (e.g., phenol) in the presence of a catalyst system where the catalyst system is activated by thermal pretreatment in a sep. reaction apparatus

L11 ANSWER 3 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2003:77802 HCAPLUS

DOCUMENT NUMBER: 138:124222

TITLE: Process and catalyst systems for the carbonylation manufacture of diaryl carbonates from phenols and carbon monoxide and dioxide

INVENTOR(S): Reisinger, Claus-Peter; Hansen, Sven Michael; Fischer, Peter

PATENT ASSIGNEE(S): Bayer A.-G., Germany; Bayer Materialscience A.-G.

SOURCE: Eur. Pat. Appl., 9 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 1279659	A2	20030129	EP 2002-15584	20020715
EP 1279659	A3	20040303		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK				
DE 10136856	A1	20030213	DE 2001-10136856	20010727
SG 103877	A1	20040526	SG 2002-4323	20020712
JP 2003096027	A2	20030403	JP 2002-211168	20020719
US 2003036663	A1	20030220	US 2002-200667	20020722
US 6852872	B2	20050208		
BR 2002002955	A	20030603	BR 2002-2955	20020725
CN 1400204	A	20030305	CN 2002-127060	20020726

PRIORITY APPLN. INFO.: DE 2001-10136856 A 20010727  
 OTHER SOURCE(S): MARPAT 138:124222

AB A process and for the carbonylation manufacture of diaryl carbonates (e.g., di-Ph carbonate) from phenols (e.g., phenol) and carbon monoxide and dioxide is conducted in the presence of a catalyst system comprising a Group VIIIB metal salt (e.g., palladium dibromide) where there are at least two metal salts (e.g., manganese trisacetylacetonate) and a base (e.g., tetrabutylammonium bromide).

L11 ANSWER 4 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2002:353405 HCAPLUS

DOCUMENT NUMBER: 136:356777

TITLE: Oxidative carbonylation process and catalyst systems for the conversion of carbon monoxide and hydroxyaromatic compounds into diaryl carbonates

INVENTOR(S): Shalyaev, Kirill Vladimirovich; Johnson, Bruce Fletcher

PATENT ASSIGNEE(S): General Electric Company, USA

SOURCE: PCT Int. Appl., 18 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002036539	A1	20020510	WO 2001-US22358	20010717
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
<del>US 6462217</del>	B1	20021008	US 2000-699829	20001030
AU 2001073504	A5	20020515	AU 2001-73504	20010717
EP 1337502	A1	20030827	EP 2001-952785	20010717
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR			
JP 2004513104	T2	20040430	JP 2002-539300	20010717
US 2003032830	A1	20030213	US 2002-163824	20020606
US 6753288	B2	20040622		

PRIORITY APPLN. INFO.: US 2000-699829 A 20001030  
 WO 2001-US22358 W 20010717

AB Organolead compds. (e.g., tetraethyllead) are useful in catalyst compns. for the oxidative carbonylation of hydroxyaromatic compds. (e.g., phenol) with oxygen and carbon monoxide into diaryl carbonates (e.g., di-Ph carbonate). The organolead compds. are employed in combination with a Group VIII metal such as palladium, or one of its compds., and a bromide or chloride such as tetraethylammonium bromide.

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 5 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2002:345990 HCAPLUS  
 DOCUMENT NUMBER: 136:356769  
 TITLE: Oxidative carbonylation process and catalysts for the production of diaryl carbonates from phenols, carbon monoxide, and oxygen  
 INVENTOR(S): Ofori, John Yaw; Pressman, Eric James; Shalyaev, Kirill Vladimirovich; Williams, Eric Douglas; Battista, Richard Anthony  
 PATENT ASSIGNEE(S): General Electric Company, USA  
 SOURCE: U.S., 13 pp., Cont.-in-part of U.S. Ser. No. 736,871. CODEN: USXXAM  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6384262	B1	20020507	US 2001-961753	20010924
WO 2002057213	A2	20020725	WO 2001-US51187	20011113
WO 2002057213	A3	20030206		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
DE 10197040	T	20031009	DE 2001-10197040	20011113
JP 2004525105	T2	20040819	JP 2002-557895	20011113
TW 574210	B	20040201	TW 2001-90129715	20011130
PRIORITY APPLN. INFO.:			US 2000-736871	A2 20001214
			US 2001-961753	A 20010924
			WO 2001-US51187	W 20011113

AB A catalytic process for the production of diaryl carbonates (e.g., di-Ph carbonate) by the oxidative carbonylation of aromatic hydroxy compds. (e.g., phenol) with carbon monoxide and oxygen is described which achieves water removal during the reaction by the steps of removing a liquid stream from an oxidative carbonylation reaction mixture in a reaction vessel, subjecting the liquid stream to reduced pressure, and returning at least a portion of the dried liquid stream to the reaction vessel. Typical oxidative carbonylation catalyst systems contain: (A) at least one Group VIII metal(s) having an atomic number of >44 or a compound the metal; (B) at least one guanidinium salt or onium salt; and (C) at least one metal co-catalyst. Process flow diagrams are provided.

REFERENCE COUNT: 48 THERE ARE 48 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 6 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2001:741565 HCAPLUS  
 DOCUMENT NUMBER: 135:289196  
 TITLE: Mixed dialkali metal salts of sulfuric acid containing cesium as polycarbonate polymerization catalysts  
 INVENTOR(S): Mccloskey, Patrick Joseph; Burnell, Timothy Brydon;

PATENT ASSIGNEE(S): Smigelski, Paul Michael, Jr.; Nisoli, Alberto  
 SOURCE: General Electric Co., USA  
 U.S., 6 pp.  
 CODEN: USXXAM  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6300460	B1	20011009	US 2000-612652	20000706
WO 2002004546	A1	20020117	WO 2001-US15497	20010514
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
EP 1301555	A1	20030416	EP 2001-935463	20010514
EP 1301555	B1	20050209		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
JP 2004502848	T2	20040129	JP 2002-509404	20010514
AT 288935	E	20050215	AT 2001-935463	20010514
TW 574256	B	20040201	TW 2001-90115352	20010626
PRIORITY APPLN. INFO.:			US 2000-612652	A 20000706
			WO 2001-US15497	W 20010514

AB The method for preparing a polycarbonate with low branched byproducts by a melt process comprises reacting a diphenol (e.g., bisphenol A) with a diaryl carbonate (e.g., di-Ph carbonate) at 100-350° in the presence of a catalyst system comprising a mixed dialkali metal salt of sulfuric acid containing at least one cesium equivalent (e.g., NaCsSO<sub>4</sub>) and a base (e.g., tetramethylammonium hydroxide).

REFERENCE COUNT: 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 7 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2001:111542 HCAPLUS

DOCUMENT NUMBER: 134:149297

TITLE: Carbonylation method and catalyst system for producing aromatic carbonates from hydroxyaromatic compounds, oxygen and carbon monoxide  
 INVENTOR(S): Patel, Ben Purushotam; Soloveichik, Grigorii Lev; Whisenhunt, Donald Wayne, Jr.; Shalyaev, Kirill Vladimirovich

PATENT ASSIGNEE(S): General Electric Company, USA

SOURCE: U.S., 7 pp.  
 CODEN: USXXAM

DOCUMENT TYPE: Patent  
 LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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US 6187942	B1	20010213	US 2000-517000	20000301
US 2001031888	A1	20011018	US 2000-729123	20001204
US 6355824	B2	20020312		
WO 2001064617	A1	20010907	WO 2001-US839	20010111
W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
EP 1263710	A1	20021211	EP 2001-955099	20010111
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
JP 2003525262	T2	20030826	JP 2001-563461	20010111
PRIORITY APPLN. INFO.: US 2000-517000 A3 20000301				
WO 2001-US839 W 20010111				

AB Aromatic hydroxy compds. (e.g., phenol) are carbonylated into diaryl carbonates (e.g., di-Ph carbonate) by contacting them with oxygen and carbon monoxide in the presence of a carbonylation catalyst system comprising an iron compound (e.g., ferrous acetate) as the primary catalyst component, and an inorg. cocatalyst (e.g., tetraethylammonium chloride). This process does not use costly platinum-group metal compound catalysts; a process flow diagram is presented.

REFERENCE COUNT: 26 THERE ARE 26 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 8 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2001:91544 HCAPLUS

DOCUMENT NUMBER: 134:149285

TITLE: Method and catalyst system for producing aromatic carbonates

INVENTOR(S): Patel, Ben Purushotam; Soloveichik, Grigorii Lev; Whisenhunt, Donald Wayne, Jr.; Shalyaev, Kirill Vladimirovich

PATENT ASSIGNEE(S): General Electric Company, USA

SOURCE: U.S., 7 pp.

CODEN: USXXAM

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6184409	B1	20010206	US 2000-516746	20000301
US 6509489	B1	20030121	US 2000-694444	20001024
WO 2001064618	A1	20010907	WO 2001-US867	20010111
W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				

EP 1261578 A1 20021204 EP 2001-901979 20010111  
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,  
 IE, SI, LT, LV, FI, RO, MK, CY, AL, TR  
 JP 2003525263 T2 20030826 JP 2001-563462 20010111  
 PRIORITY APPLN. INFO.: US 2000-516746 A3 20000301  
 WO 2001-US867 W 20010111

AB The method comprises the step of contacting  $\geq 1$  aromatic hydroxy compound with oxygen and CO in the presence of a carbonylation catalyst system having an effective amount of a nickel source as the primary catalyst component and optionally  $\geq 1$  inorg. co-catalyst, as well as a halide composition and/or a base in the absence of a Group VIII B metal source. A process flow diagram is presented.

REFERENCE COUNT: 26 THERE ARE 26 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 9 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2001:45205 HCAPLUS

DOCUMENT NUMBER: 134:87919

TITLE: Carbonylation process and catalyst system for producing diaryl carbonates from the reaction of carbon monoxide and oxygen with hydroxyaromatic compounds

INVENTOR(S): Patel, Ben Purushotam; Soloveichik, Grigorii Lev; Whisenhunt, Donald Wayne, Jr.; Shalyaev, Kirill Vladimirovich

PATENT ASSIGNEE(S): General Electric Company, USA

SOURCE: U.S., 6 pp.  
 CODEN: USXXAM

DOCUMENT TYPE: Patent  
 LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6175033	B1	20010116	US 2000-510381	20000222
WO 2001062702	A1	20010830	WO 2000-US29285	20001024
W. AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
EP 1261576	A1	20021204	EP 2000-973807	20001024
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL				
US 6380418	B1	20020430	US 2000-721682	20001127
PRIORITY APPLN. INFO.: US 2000-510381 A 20000222				
WO 2000-US29285 W 20001024				

AB A method of carbonylating aromatic hydroxy compds. into a diaryl carbonate (e.g., di-Ph carbonate) comprises reacting at least one aromatic hydroxy compound (e.g., phenol) with oxygen and carbon monoxide in the presence of a carbonylation catalyst system comprising an effective amount of a manganese source [e.g., manganese(II) acetylacetonate] as a primary catalyst component in the absence of a Group VIIIB metal source, and, optionally in the presence of of a catalytic amount of an inorg. cocatalyst [e.g., lead(II) oxide] as well as a halide composition

(e.g., tetraethylammonium bromide), and/or a base. A process flow diagram is presented.

REFERENCE COUNT: 26 THERE ARE 26 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 10 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 2000:441754 HCAPLUS

DOCUMENT NUMBER: 133:75637

TITLE: Method for processing reaction mixtures containing diaryl carbonate

INVENTOR(S): Hesse, Carsten; Jansen, Ursula; Rechner, Johann; Reisinger, Claus-Peter; Eek, Rob; Hallenberger, Kaspar; Friedrich, Martin

PATENT ASSIGNEE(S): Bayer Aktiengesellschaft, Germany

SOURCE: PCT Int. Appl., 21 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000037419	A1	20000629	WO 1999-EP9697	19991209
W:				
AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW:				
GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
DE 19859295	A1	20000629	DE 1998-19859295	19981222
BR 9916464	A	20010925	BR 1999-16464	19991209
EP 1140782	A1	20011010	EP 1999-963455	19991209
EP 1140782	B1	20030903		
R:				
AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
JP 2002533315	T2	20021008	JP 2000-589491	19991209
AT 248800	E	20030915	AT 1999-963455	19991209
ES 2207320	T3	20040516	ES 1999-963455	19991209
US 6605191	B1	20030812	US 2001-868298	20010615
PRIORITY APPLN. INFO.:			DE 1998-19859295	A 19981222
			WO 1999-EP9697	W 19991209

OTHER SOURCE(S): MARPAT 133:75637

AB The invention relates to a method for processing reaction mixts. containing diaryl carbonate, an aromatic hydroxy compound, water, a base, a quaternary salt and other catalyst components, said reaction mixts. being obtained in the production of diaryl carbonates by direct carbonylation of aromatic hydroxy compds. The reaction mixture is distilled at 80-160°/1-100 mbar and 1 theor. distillation plate to give a gas phase containing a diaryl carbonate, an aromatic hydroxy compound, and water and a liquid phase containing a diaryl carbonate, aromatic hydroxy compound, base, a quaternary salt, and other catalyst components for recycling to the carbonylation. This process is performed in film evaporators and provides for the recycling with a min. loss of activity of the catalyst system.



REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 11 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN  
 ACCESSION NUMBER: 2000:157747 HCAPLUS  
 DOCUMENT NUMBER: 132:182331  
 TITLE: Continuous oxidative carbonylation process  
 and catalyst system for the  
 manufacture of diaryl carbonates  
 from hydroxyaromatic compounds and oxygen and carbon  
 monoxide  
 INVENTOR(S): Moreno, Phillip  
 PATENT ASSIGNEE(S): General Electric Company, USA  
 SOURCE: U.S., 8 pp.  
 CODEN: USXXAM  
 DOCUMENT TYPE: Patent  
 LANGUAGE: English  
 FAMILY ACC. NUM. COUNT: 1  
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6034262	A	20000307	US 1998-218651	19981222
WO 2000037417	A1	20000629	WO 1999-US24528	19991020
W: CN, JP, SG				
RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
EP 1140777	A1	20011010	EP 1999-955073	19991020
EP 1140777	B1	20040526		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI				
JP 2002533313	T2	20021008	JP 2000-589489	19991020
AT 267794	E	20040615	AT 1999-955073	19991020
PRIORITY APPLN. INFO.:			US 1998-218651	A 19981222
			WO 1999-US24528	W 19991020

OTHER SOURCE(S): MARPAT 132:182331

AB Diaryl carbonates (e.g., di-Ph carbonate) are manufactured in a continuous process by contacting at least one hydroxyarom. compound (e.g., phenol) with oxygen and carbon monoxide in the presence of catalyst system comprising a Group VIIIB metal catalyst [e.g., Pd(acac)<sub>2</sub>], an inorg. co-catalyst (e.g., PbO), an optional organic catalyst, and at least one halide source (e.g., hexaethylguanidinium bromide), in which one provides a first solution comprising at least one first catalyst system component in a first tank, a second solution comprising at least one second catalyst system component in a second tank, and feeding the first and second solns. sep. into a reactor.

REFERENCE COUNT: 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 12 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN  
 ACCESSION NUMBER: 1999:285937 HCAPLUS  
 DOCUMENT NUMBER: 130:282484  
 TITLE: Carbonylation process and  
 hexaalkylguanidinium halide-containing  
 catalyst system for preparing  
 diaryl carbonates from  
 hydroxyaromatic compounds  
 INVENTOR(S): Pressman, Eric James; Shafer, Sheldon Jay

PATENT ASSIGNEE(S): General Electric Company, USA  
SOURCE: U.S., 3 pp., Cont.-in-part of U.S. Ser. No. 40,264.  
CODEN: USXXAM  
DOCUMENT TYPE: Patent  
LANGUAGE: English  
FAMILY ACC. NUM. COUNT: 2  
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5898079	A	19990427	US 1997-929000	19970912
US 5898080	A	19990427	US 1997-40264	19970827
JP 10306065	A2	19981117	JP 1998-21365	19980203
EP 858991	A1	19980819	EP 1998-300911	19980209
EP 858991	B1	20020116		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
ES 2169894	T3	20020716	ES 1998-300911	19980209
CN 1211564	A	19990324	CN 1998-104450	19980213
PRIORITY APPLN. INFO.:			US 1997-40264	A2 19970808
			US 1997-40300P	P 19970213
			US 1997-40264P	P 19970827
			US 1997-929000	A 19970912

AB Diaryl carbonates, useful as polycarbonate monomers (no data), are prepared in high yield and selectivity by contacting  $\geq 1$  hydroxyarom. compound with oxygen and carbon monoxide in the presence of catalyst system comprising palladium or its compds., a lead compound (i.e., an inorg. cocatalyst), and  $\geq 1$  of a hexaalkylguanidinium bromide and/or chloride, where the use of the hexaalkylguanidinium salt causes an increase in the yield of the diaryl carbonate without a decrease in selectivity. Thus, phenol, hexaethylguanidinium bromide, lead(II) oxide, and palladium(II) 2,4-pentanedioate were added to a reactor, the reactor pressurized with 4150 kg/m<sup>2</sup> of carbon monoxide and 2075 kg/m<sup>2</sup> of air, heated at 100°, and a mixture of carbon monoxide and air introduced over 6 h, producing a di-Ph carbonate yield of 11.4% and a palladium turnover number of 11,534, while a control carbonylation using tetra-n-butylammonium bromide instead of hexaethylguanidinium bromide showed a yield of 10.0% and a palladium turnover number of 8970.

REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 13 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1998:653707 HCAPLUS

DOCUMENT NUMBER: 129:245657

TITLE: Carbonylation process and stable  $\beta$ -diketone salt catalysts for preparing diaryl carbonates from hydroxyaromatic compounds

INVENTOR(S): Pressman, Eric James; Shafer, Sheldon Jay

PATENT ASSIGNEE(S): GENERAL ELECTRIC COMPANY, USA

SOURCE: Eur. Pat. Appl., 6 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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EP 867428 A1 19980930 EP 1998-302049 19980318  
EP 867428 B1 20040303  
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, MC, PT, IE,  
SI, LT, LV, FI, RO  
US 5908952 A 19990601 US 1997-823784 19970324  
SG 77629 A1 20010116 SG 1998-550 19980311  
JP 10330325 A2 19981215 JP 1998-64848 19980316  
CN 1194261 A 19980930 CN 1998-105858 19980324  
PRIORITY APPLN. INFO.: US 1997-823784 A 19970324

AB Hydroxyarom. compds. (e.g., phenol) are carbonylated to diaryl carbonates (e.g., di-Ph carbonate) by reaction with oxygen and carbon monoxide in the presence of a catalyst system which comprises a Group VIII metal salt of an aliphatic  $\beta$ -diketone [e.g., palladium(II) 2,4-pentanedionate], and, optionally, an inorg. cocatalyst, an organic cocatalyst (e.g., 2,2':6',2''-terpyridine), and a bromide or chloride source (e.g., hexethylguanidinium bromide). The use of the  $\beta$ -diketone salt confers long shelf life under normal storage conditions (i.e., nonpptn. of the Pd), high activity upon recycle, and capability of carbonylation at  $<100^\circ$ .

REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 14 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1998:568638 HCAPLUS

DOCUMENT NUMBER: 129:176095

TITLE: Carbonylation method for preparing diaryl carbonate monomers from hydroxyaromatic compounds using catalyst systems containing hexaalkylguanidinium chlorides or bromides

INVENTOR(S): Pressman, Eric James; Shafer, Sheldon Jay

PATENT ASSIGNEE(S): General Electric Co., USA

SOURCE: Eur. Pat. Appl., 5 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 858991	A1	19980819	EP 1998-300911	19980209
EP 858991	B1	20020116		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, SI, LT, LV, FI, RO				
US 5898080	A	19990427	US 1997-40264	19970827
US 5898079	A	19990427	US 1997-929000	19970912
CN 1194260	A	19980930	CN 1998-105856	19980324

PRIORITY APPLN. INFO.:  
US 1997-40300P P 19970213  
US 1997-40264 P 19970827  
US 1997-929000 A 19970912

AB Hydroxyarom. compds. (e.g., phenol) are converted into diaryl carbonates (e.g., di-Ph carbonate) in high yield and selectivity by reaction with oxygen and carbon monoxide in the presence of a catalyst system comprising a Group VIIIB metal or compound [e.g., palladium(II) acetate], an inorg. cocatalyst [e.g., a cobalt(II) complex with bis[3-(salicylal amino)propyl]methylamine], an organic cocatalyst (e.g., 2,2':6',2''-terpyridine), and a hexaalkylguanidinium bromide or chloride (e.g., hexaethylguanidinium bromide). The use of the

hexaalkylguanidinium bromide or chloride causes an increase in the yield of the diaryl carbonate product without a decrease in its selectivity of formation.

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 15 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1997:682236 HCAPLUS

DOCUMENT NUMBER: 127:307214

TITLE: Process for the continuous production of diaryl carbonates by the oxidative carbonylation of aromatic hydroxy compounds

INVENTOR(S): Buysch, Hans-Josef; Hesse, Carsten; Rechner, Johann

PATENT ASSIGNEE(S): Bayer A.-G., Germany

SOURCE: Eur. Pat. Appl., 14 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 801051	A1	19971015	EP 1997-105177	19970327
EP 801051	B1	20020327		
R: BE, DE, ES, FR, GB, IT, NL				
DE 19614062	A1	19971016	DE 1996-19614062	19960409
US 5712406	A	19980127	US 1997-794435	19970205
ES 2174141	T3	20021101	ES 1997-105177	19970327
US 5821377	A	19981013	US 1997-825603	19970401
JP 10036323	A2	19980210	JP 1997-100764	19970404
CN 1168879	A	19971231	CN 1997-110535	19970409
CN 1169422	A	19980107	CN 1997-111037	19970516
PRIORITY APPLN. INFO.:			DE 1996-19614062	A 19960409

OTHER SOURCE(S): MARPAT 127:307214

AB Diaryl carbonates (e.g., di-Ph carbonate) are prepared by the oxidative carbonylation of aromatic hydroxy compds. (e.g., PhOH) in the presence of a catalyst system comprising a platinum-group metal, a co-catalyst, and a quaternary salt or base. Process flow diagrams are presented.

L11 ANSWER 16 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1995:655204 HCAPLUS

DOCUMENT NUMBER: 123:55489

TITLE: Preparation of diaryl carbonates

INVENTOR(S): Buysch, Hans-Josef; Dohm, Joachim; Hesse, Carsten; Rechner, Johann; Kaufmann, Dieter

PATENT ASSIGNEE(S): Bayer A.-G., Germany

SOURCE: Eur. Pat. Appl., 11 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 654461	A1	19950524	EP 1994-117665	19941109
EP 654461	B1	19971203		

R: BE, CH, DE, ES, FR, GB, IT, LI, NL

DE 4339697	A1	19950524	DE 1993-4339697	19931122
DE 4341990	A1	19950614	DE 1993-4341990	19931209
ES 2110683	T3	19980216	ES 1994-117665	19941109
US 5502232	A	19960326	US 1994-339613	19941115
JP 07188116	A2	19950725	JP 1994-305701	19941116
CA 2135656	AA	19950523	CA 1994-2135656	19941118
CN 1107833	A	19950906	CN 1994-118957	19941122
CN 1054836	B	20000726		

PRIORITY APPLN. INFO.:

DE 1993-4339697	A	19931122
DE 1993-4341990	A	19931209

OTHER SOURCE(S): CASREACT 123:55489

AB (RO)2CO [R = (un)substituted aryl] were prepared in a process in which an aromatic hydroxy compound is condensed with CO in the presence of O, a drying agent, and a catalyst system comprising a noble metal, a base, a quaternary salt, and a cocatalyst, the metal catalyst being activated by CO pretreatment in the presence of the quaternary salt and, optionally, the cocatalyst. Thus, PdBr<sub>2</sub> and Bu<sub>4</sub>NBr in PhOH containing 750ppm H<sub>2</sub>O at 55° were treated with CO after which Zeolite A, Mn(acac)<sub>2</sub>, and pentamethylpiperidine were added and an air/CO (1:1) mixture introduced for 6h to give a mixture comprising 1.5% (PhO)2CO.

L11 ANSWER 17 OF 17 HCAPLUS COPYRIGHT 2006 ACS on STN

ACCESSION NUMBER: 1994:191360 HCAPLUS

DOCUMENT NUMBER: 120:191360

TITLE: Preparation of aromatic carbonic acid esters

INVENTOR(S): Iwane, Hiroshi; Myagi, Hidekazu; Imada, Satoshi; Seo, Shoichi; Yoneyama, Takahiro

PATENT ASSIGNEE(S): Mitsubishi Petrochemical Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 4 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 06009505	A2	19940118	JP 1992-161180	19920619
JP 3128329	B2	20010129		

PRIORITY APPLN. INFO.:

JP 1992-161180	19920619
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OTHER SOURCE(S): CASREACT 120:191360

AB Aromatic carbonic acid esters are prepared by reaction of aromatic hydroxy compds., CO, and O in the presence of catalysts (A)  $\geq 1$  Pd and Pd compds., (B)  $\geq 1$  Ce(III) and Ce(IV) compds., (C)  $\geq 1$  quaternary ammonium and phosphonium salts, and (D)  $\geq 1$  quinone and its reduced products, aromatic diols. This process suppresses the formation of oxidative dimerization and trimerization byproducts such as p-phenoxyphenol which has a b.p. close to that of (PhO)2CO and is difficult to sep., and gives the desired products in high yields. Thus, 7.8 g phenol, Pd(OAc)<sub>2</sub> 2.4, Ce(OAc)<sub>3</sub>·H<sub>2</sub>O 3.5, Bu<sub>4</sub>NBr 202, and hydroquinone 34 mg were charged in a Hastelloy autoclave; after flushing the system with CO, 60 atom CO and 30 atom dry air were introduced; and the mixture was allowed to react at 120° for 1 h to give (PhO)2CO 3.7, Ph salicylate 0.12, and p-phenoxyphenol 0.039% (1.0% selectivity). Diaryl carbonates, particularly (PhO)2CO, are useful as intermediates for polycarbonates.

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COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
93.38	93.59

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE	TOTAL
ENTRY	SESSION
-17.25	-17.25

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STN INTERNATIONAL LOGOFF AT 10:41:52 ON 26 SEP 2006